

BARNES & THORNBURG LLP

11 South Meridian Street
Indianapolis, Indiana 46204
(317) 236-1313
(317) 231-7433 Fax

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Group: 1626

Confirmation No.: 9842

Application No.: 10/725,191

Invention: **PHOSPHORAMIDE
COMPOUNDS**

Applicant: Richard F. Borch et al.

Filed: December 1, 2003

Attorney Docket: 3220-200986

Examiner: Golam M. Shameem

I hereby certify that this correspondence is being
filed electronically in the United States Patent and
Trademark Office

on March 23, 2007


(Signature)

Garla L. Twyman
(Printed Name)

SUBMISSION MADE WITH AN RCE:

Supplemental Information Disclosure Statement

Mail Stop RCE

Commissioner for Patents

P. O. Box 1450

Alexandria, Virginia 22313-1450

Sir:

Applicants have concurrently filed a request for continued examination (RCE) under 37 C.F.R. § 1.114 in a separate paper. In association with that RCE, Applicants hereby request entry and consideration of the instant Submission, which Submission includes a Supplemental Information Disclosure Statement (IDS), Form 1449, and all required copies of the references cited therein.

This IDS is filed in the application identified above pursuant to 37 C.F.R. § 1.56. No representation is intended that a complete search has been made of the prior art or that no better art references than the references cited in the IDS are available. The filing of this IDS shall not be construed to be an admission that the information cited in the IDS is, or

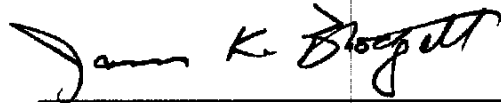
1.98(a)(2)(ii), copies of the cited U.S. patents are not provided herewith. Copies of the cited foreign patent documents and other references (non-patent) have been provided.

The cited references are not believed to disclose or suggest the invention recited in the claims of the above-identified application. It is therefore believed that the claimed invention is patentably distinguishable over the cited references.

Applicants have authorized, concurrent to filing the instant IDS, Barnes & Thornburg LLP Deposit Account 10-0435 to be debited in the amount of \$575.00, which amount is the sum of the \$180.00 fee set forth under 37 C.F.R. 1.17(p) for filing an Information Disclosure Statement, and the \$395.00 fee set forth under 37 C.F.R. § 1.17(e) for requesting continued examination under small-entity status.

Applicants do not believe that any additional fees are due for filing the instant Information Disclosure Statement. However, the Commissioner is hereby authorized to charge any additional fees that may be required, or credit any overpayment, to Barnes & Thornburg LLP Deposit Account No. 10-0435, with reference to our matter number 3220-200986.

Respectfully submitted,
BARNES & THORNBURG LLP



James K. Blodgett
Registration No. 48,480
Agent for Applicants

JKB:glt
Indianapolis, Indiana 46204
317-231-7401

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE INFORMATION DISCLOSURE STATEMENT				ATTY. DOCKET NO.: 3220-200986 APPLICANT(S): Borch et al. FILING DATE: Dec. 1, 2003		SERIAL NO.: 10/725,191 GROUP: 1626	
U.S. PATENT DOCUMENTS							
*Examiner Initial		Document Number	Date	Name	Class	Subclass	Filing Date if appropriate
	AA	5,233,031					
	AB	6,903,081					
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		Document Number	Date	Country	Class	Subclass	Translation Yes No
	AL	WO 93/06120	April 1, 1993	PCT			
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	AN						
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OTHER REFERENCES (Including Author, Title, Date, Pertinent Pages, Etc.)							
	AR	"The Synthesis of O-Monosaccharidyl-Methoxycarbonyl-Phosphonamidaes by Arbuzov Reaction," Chen et al., Chinese Chemical Letters, vol. 6, No. 1, pp. 23-26, 1995					
	AS	"Design and Synthesis of Lipophilic Phosphoramidate d4T-MP Prodrugs Expressing High Potency Against HIV in Cell Culture: Structural Determinants for <i>in vitro</i> Activity and QSAR," Siseiqli, et al., J. Med. Chem. 1999, 42, 4122-4128					
	AT	"Phosphoramidates as Potent Prodrugs of Anti-HIV Nucleotides: Studies in the Amino Region," McGuigan et al., Antiviral Chemistry & Chemotherapy (1996) 7(1), 31-36					
	AU	"Design and Synthesis of Novel Nucleotide Prodrugs," Meyers et al., American Association for Cancer Research 2000 (Abstract Proof Page), Abstract #100710, dated December 2, 1999					
	AV	"Protein Tyrosine Kinases and Cancer," Kolibaba et al., "Biochimica et Biophysica Acta 1333 (1997) F217-F248					
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	AZ	"Oncogenes and Signal Transduction," Cantley et al., Cell, Vol. 64, January 25, 1991, 281-302					
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*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.							

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	BL						
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	BR	"Recognition and Specificity in Protein Tyrosine Kinase-Mediated Signalling," Songyang et al., TIBS 20, November 1995, 470-475
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	BT	"Peptide Inhibitors of src SH3-SH2-Phosphoprotein Interactions," Gilmer et al., The Journal of Biological Chemistry, Vol. 269, No. 50, December 16, 1994, 31711-31719
	BU	"Structure-Based Design of a Novel Series of Nonpeptide Ligands That Bind to the pp60 ^{src} SH2 Domain," Lunney et al., J. Am. Chem. Soc. 1997, 119, 12471-12476
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	BX	"Nonhydrolyzable Phosphotyrosyl Mimetics for the Preparation of Phosphatase-resistant SH2 Domain Inhibitors," Burke, Jr., et al., Biochemistry, 1994, 33, 6490-6494
	BY	L-O-(2-Malonyl)tyrosine: A New Phosphotyrosyl Mimetic for the Preparation of Src Homology 2 Domain Inhibitory Peptides," Ye et al., J. Med. Chem. 1995, 38, 4270-7275
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	CL						
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	CR	"4'-O-[2-(2-Fluoromalonyl)]-L-Tyrosine: A Phosphotyrosyl Mimic for the Preparation of Signal Transduction Inhibitory Peptides," Burke Jr., et al., J. Med. Chem. 1996, 39, 1021-1027					
	CS	"Phosphotyrosine-Containing Dipeptides as High-Affinity Ligands for the p56 ^{lck} SH2 Domain," Llinas-Brunet et al., J. Med. Chem. 1999, 42, 722-729					
	CT	"Acquisition of High-Affinity, SH2-Targeted Ligands via a Spatially Focused Library," Lee et al., J. Med. Chem. 1999, 42, 784-787					
	CU	"Ligands for the Tyrosine Kinase p56 ^{lck} SH2 Domain: Discovery of Potent Dipeptide Derivatives with Monocharged, Nonhydrolyzable Phosphate Replacements," Beaulieu et al., J. Med. Chem. 1999, 42, 1757-1766					
	CV	"Synthesis and Biological Evaluation of 5-Fluoro-2'-Deoxyuridine Phosphoramidate Analogs," Fries et al., J. Med. Chem., 1995, 38, 2672-2680					
	CW	"Synthesis and Biological Activity of Novel 5-Fluoro-2'-Deoxyuridine Phosphoramidate Prodrugs," Meyers et al., J. Med. Chem., 2000, 43, 4313-4318					
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	CY	Gibbs et al., Current Medicinal Chemistry, Vol. 8, No. 12, October 2001, pp. 1437-1465					
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	DR	Weidong et al., American Chemical Society. Abstracts of Papers at the National Meeting, ACS, Washington, D.C., Vol. 228, Part 1, 26 August 2004, p. U930
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